

The new substrate is synthetically easily accessible

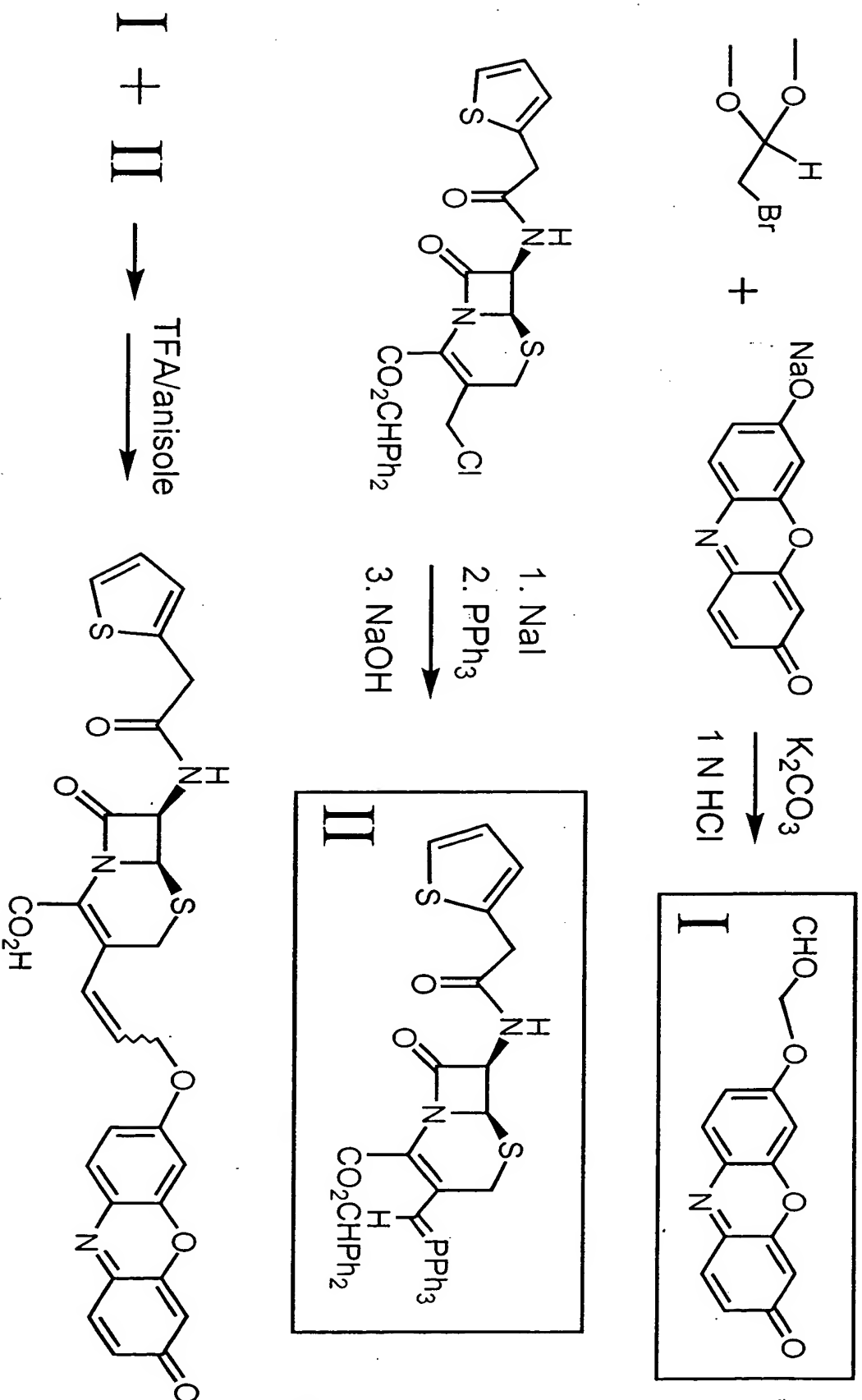


FIG. 1

Enzymatic fragmentation can take place to the new substrate

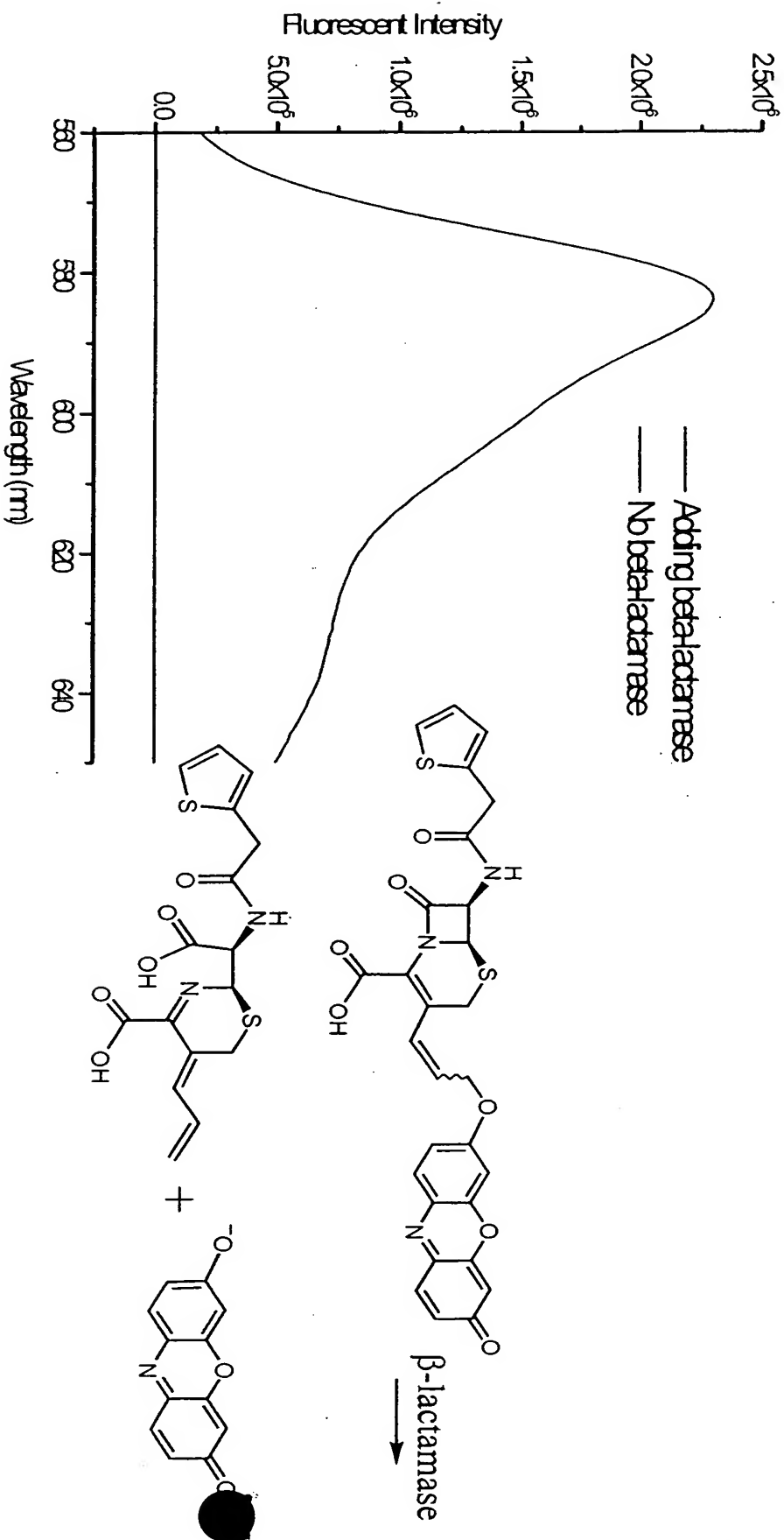


FIG. 2

Synthesis of RECTO

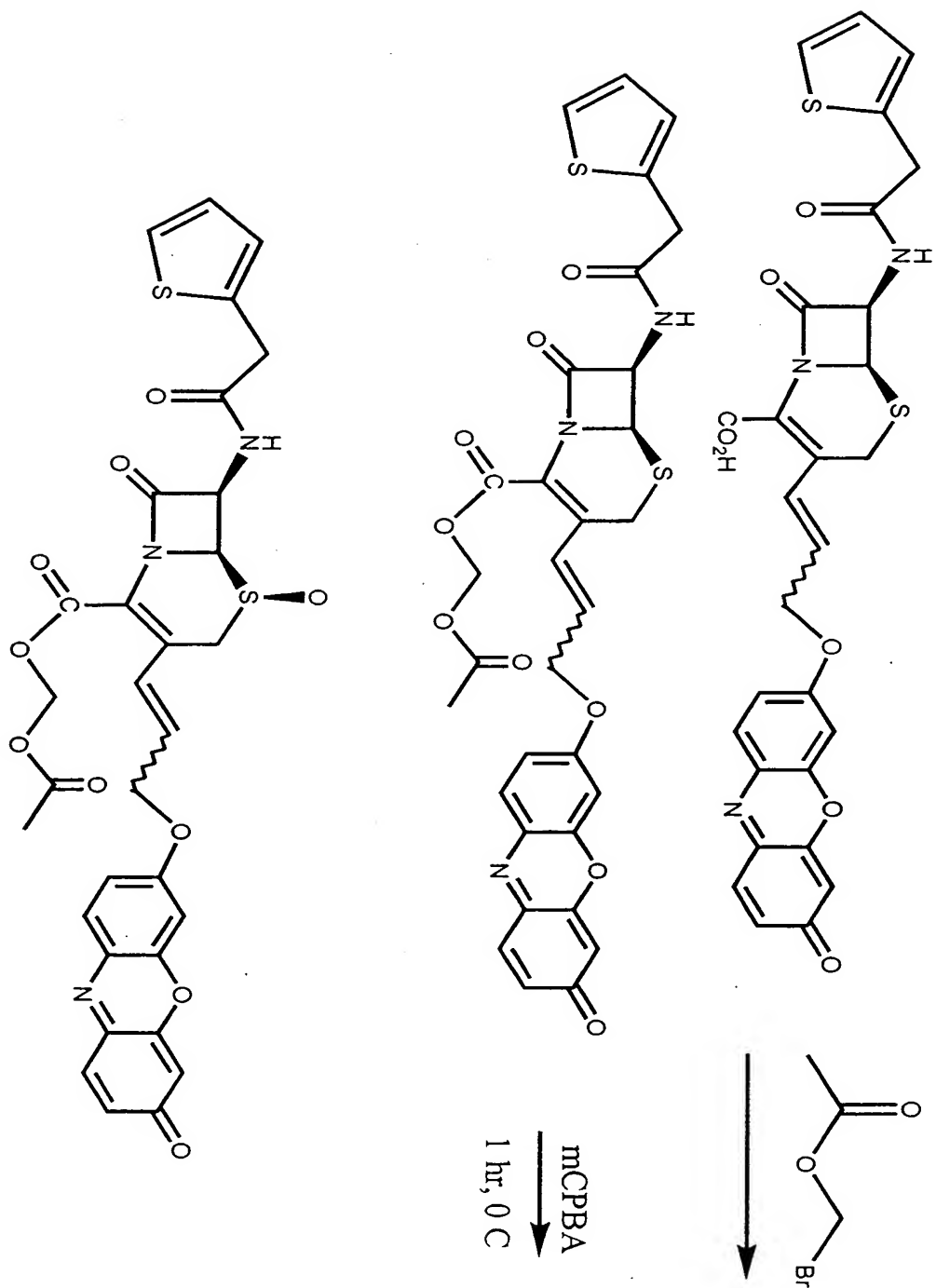
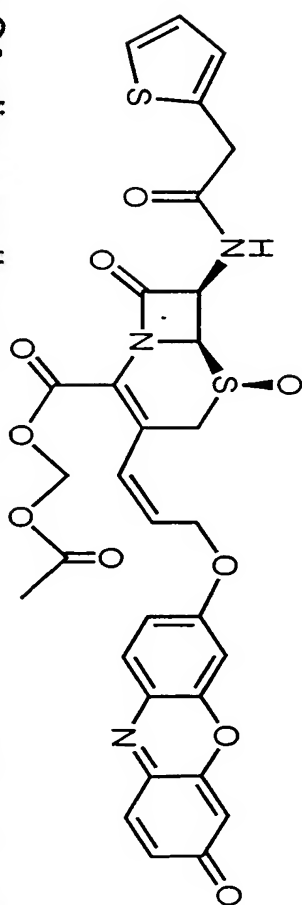


FIG. 3





Increased resorufin deposition in β -lactamase-transfected vs. wild type cells



BLA-transfected C6 glioma cells



WT C6 glioma

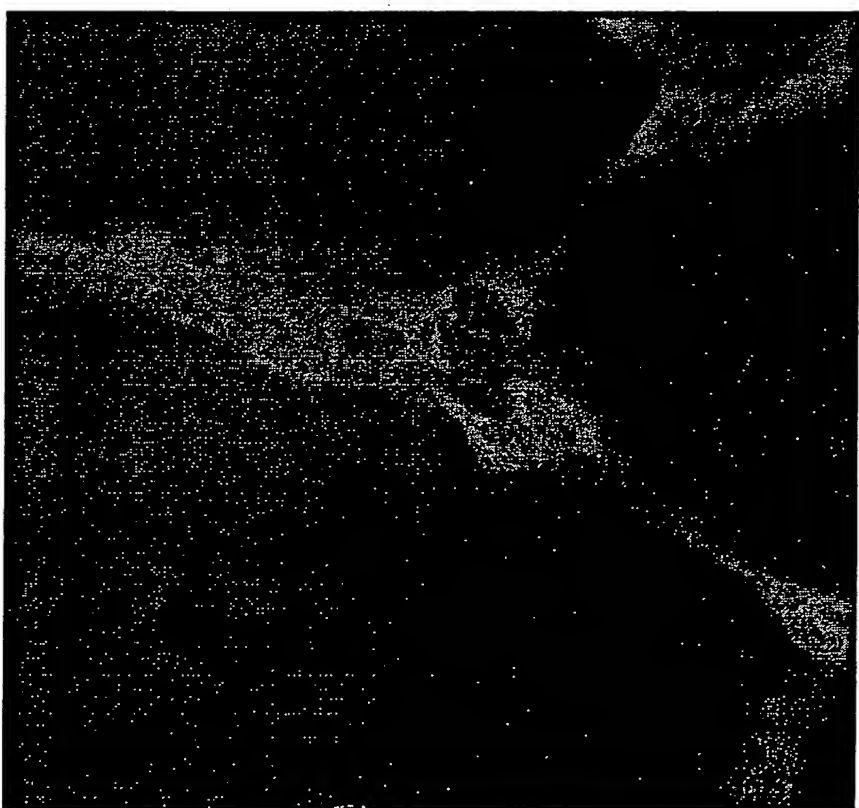
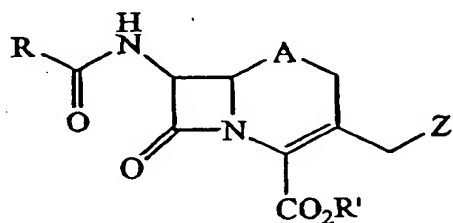


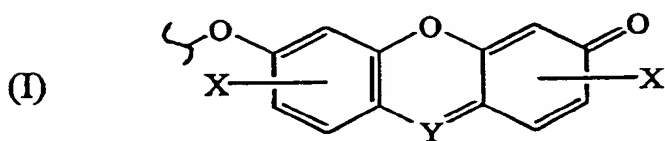
FIG. 6

cephalosporin-phenol ethers that we wish to claim:



Preferred R = benzyl, 2-thienylmethyl, or cyanomethyl; A = S or SO; R' = H or physiologically acceptable salts or ester groups.

where Z can be:



where X = H, F, Cl, Br, CO₂R';
Y = N, CH, C-CN, C-CF₃

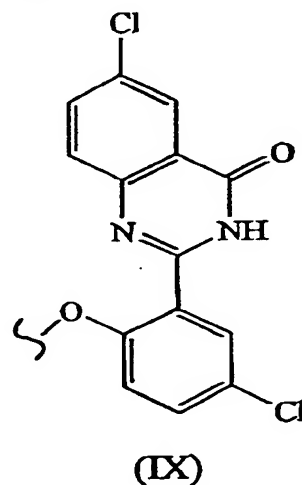
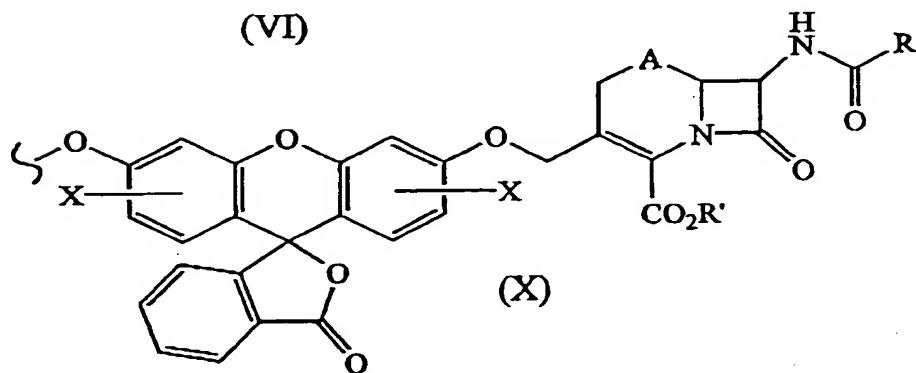
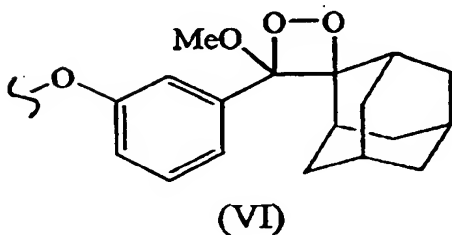
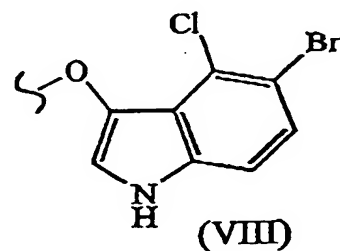
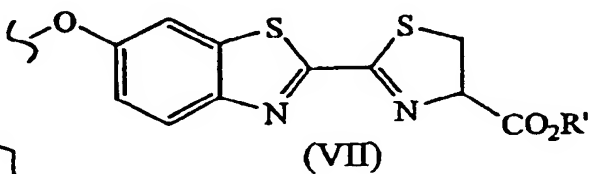
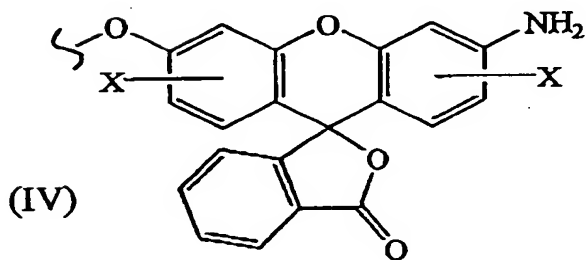
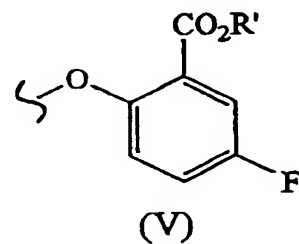
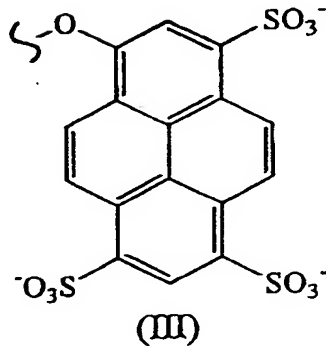
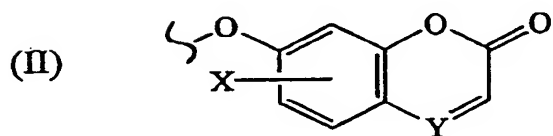
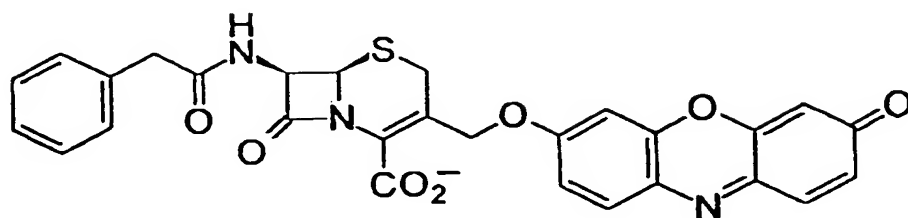


FIG. 7

Resorufin-cephalosporin cleaved by β -lactamase



β -lactamase

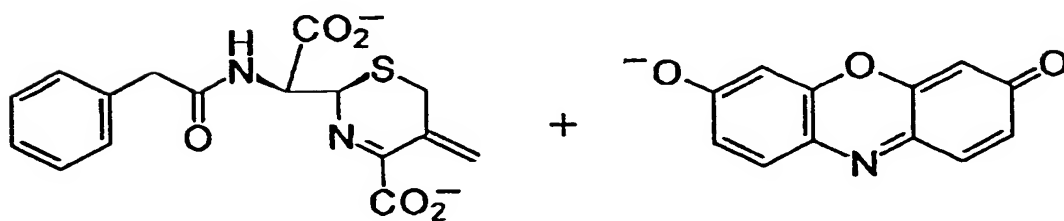


FIG. 8

Absorption spectra of resorufin-cephalosporin before and after β -lactamase treatment

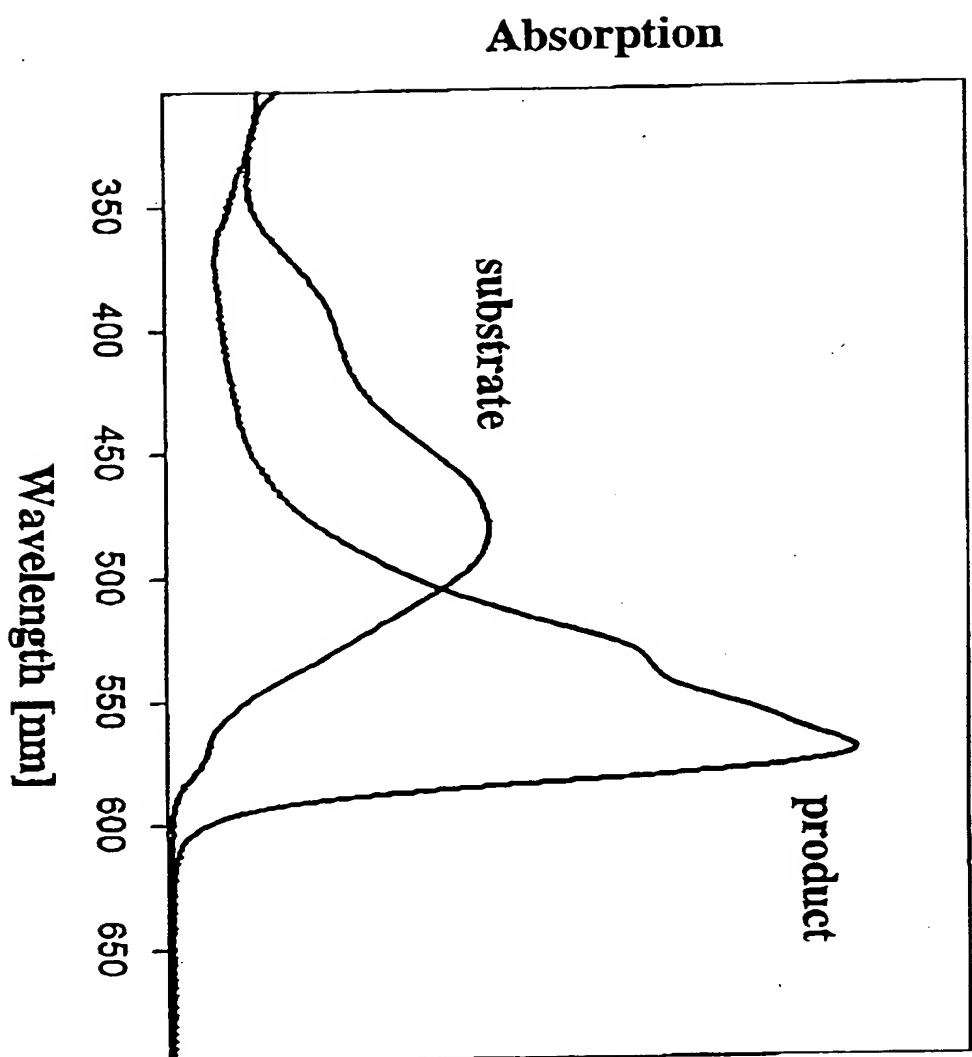


FIG. 9

Fluorescence emission of resorufin-cephalosporin before and after β -lactamase treatment (excitation at 570 nm)

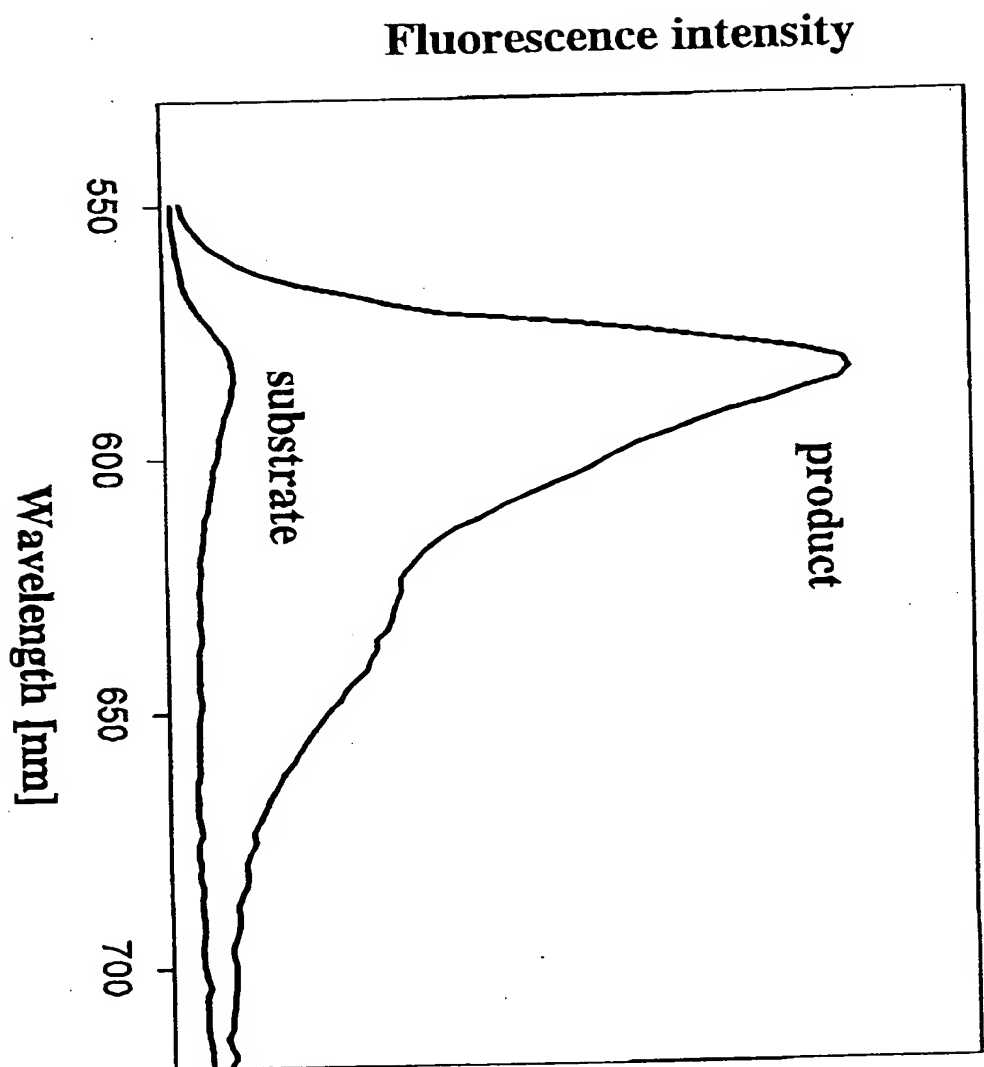


FIG. 10